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NEWS 19 FEB 16 STN User Update to be held in conjunction with the 229th ACS  
National Meeting on March 13, 2005  
  
NEWS EXPRESS JANUARY 10 CURRENT WINDOWS VERSION IS V7.01a, CURRENT  
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\* \* \* \* \* STN Columbus \* \* \* \* \*

FILE 'HOME' ENTERED AT 09:58:56 ON 19 FEB 2005

=> file caplus  
COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
0.21	0.21

FULL ESTIMATED COST

FILE 'CAPLUS' ENTERED AT 09:59:03 ON 19 FEB 2005

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FILE COVERS 1907 - 19 Feb 2005 VOL 142 ISS 9

FILE LAST UPDATED: 18 Feb 2005 (20050218/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> HCV (s) replicon

8738 HCV

17 HCVS

8742 HCV

(HCV OR HCVS)

2986 REPLICON

1500 REPLICONS

3685 REPLICON

(REPLICON OR REPLICONS)

L1 239 HCV (S) REPLICON

=> lambda (w) phage

171690 LAMBDA

66 LAMBDA

171703 LAMBDA

(LAMBDA OR LAMBDA)

45082 PHAGE

7309 PHAGES

46708 PHAGE

(PHAGE OR PHAGES)

L2 2782 LAMBDA (W) PHAGE

=> L1 and L2

L3 0 L1 AND L2

=> phage

45082 PHAGE

7309 PHAGES

L4 46708 PHAGE

(PHAGE OR PHAGES)

=> L1 and L4

L5 1 L1 AND L4

=> D L5 IBIB ABS

L5 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2004:1010148 CAPLUS

DOCUMENT NUMBER: 142:130666

TITLE: Protein kinase C-related kinase 2 regulates hepatitis C virus RNA polymerase function by phosphorylation

AUTHOR(S): Kim, Seong-Jun; Kim, Jung-Hee; Kim, Yeon-Gu; Lim, Ho-Soo; Oh, Jong-Won

CORPORATE SOURCE: Department of Biotechnology, Yonsei University, Seoul, 120-749, S. Korea

SOURCE: Journal of Biological Chemistry (2004), 279(48), 50031-50041

CODEN: JBCHA3; ISSN: 0021-9258

PUBLISHER: American Society for Biochemistry and Molecular Biology

DOCUMENT TYPE: Journal

LANGUAGE: English

AB The hepatitis C virus (HCV) NS5B protein is the viral RNA-dependent RNA polymerase required for replication of the HCV RNA genome. We have identified a peptide that most closely resembles a short region of the protein kinase C-related kinase 2 (PRK2) by screening of a random 12-mer peptide library displayed on the surface of the M13 bacteriophage with NS5B proteins immobilized on microwell plates. Competitive phage ELISA with a synthetic peptide showed that the phage clone displaying this peptide could bind HCV RNA polymerase with a high affinity. Coimmunopptn. and colocalization studies demonstrated in vivo interaction of NS5B with PRK2. In vitro kinase assays demonstrated that PRK2 specifically phosphorylates NS5B by interaction with the N-terminal finger domain of NS5B (amino acids 1-187). Consistent with the in vitro NS5B-phosphorylating activity of PRK2, we detected the phosphorylated form of NS5B by metabolic cell labeling. Furthermore, HCV NS5B immunopptd. from HCV subgenomic replicon cells was specifically recognized by an antiphosphoserine antibody. Knock-down of the endogenous PRK2 expression using a PRK2-specific small interfering RNA inhibited HCV RNA replication. In contrast, PRK2 overexpression, which was accompanied by an increase of in the level of its active form, dramatically enhanced HCV RNA replication. Altogether, our results indicate that HCV RNA replication is regulated by NS5B phosphorylation by PRK2.

REFERENCE COUNT: 74 THERE ARE 74 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT